FDA 510(k) Premarket Notification

K183071 VLIFT-s Vertebral Body Replacement System

CLEARED

Jan 10, 2019
66 days to decision

K183071 · Product code: **PLR** · Orthopedic
Source: https://www.510kdatabase.net/k183071/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Spinal Vertebral Body Replacement Device - Cervical (PLR)
Date received	Nov 5, 2018
Decision date	Jan 10, 2019
Days to decision	66 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Stryker Spine**
Location	Allendale, NJ, US
Contact	Renee Norby
510(k) history	74 submissions · 73 cleared · 2004-2026